

10/773,002

NEWS 3 FEB 28 PATDPAFULL - New display fields provide for legal status
data from INPADOC
NEWS 4 FEB 28 BABS - Current-awareness alerts (SDIs) available
NEWS 5 MAR 02 GBFULL: New full-text patent database on STN
NEWS 6 MAR 03 REGISTRY/ZREGISTRY - Sequence annotations enhanced
NEWS 7 MAR 03 MEDLINE file segment of TOXCENTER reloaded
NEWS 8 MAR 22 KOREAPAT now updated monthly; patent information enhanced
NEWS 9 MAR 22 Original IDE display format returns to REGISTRY/ZREGISTRY
NEWS 10 MAR 22 PATDPASPC - New patent database available
NEWS 11 MAR 22 REGISTRY/ZREGISTRY enhanced with experimental property tags
NEWS 12 APR 04 EPFULL enhanced with additional patent information and new
fields
NEWS 13 APR 04 EMBASE - Database reloaded and enhanced
NEWS 14 APR 18 New CAS Information Use Policies available online
NEWS 15 APR 25 Patent searching, including current-awareness alerts (SDIs),
based on application date in CA/CAPLUS and USPATFULL/USPAT2
may be affected by a change in filing date for U.S.
applications.
NEWS 16 APR 28 Improved searching of U.S. Patent Classifications for
U.S. patent records in CA/CAPLUS
NEWS 17 MAY 23 GBFULL enhanced with patent drawing images
NEWS 18 MAY 23 REGISTRY has been enhanced with source information from
CHEMCATS
NEWS 19 JUN 06 The Analysis Edition of STN Express with Discover!
(Version 8.0 for Windows) now available
NEWS 20 JUN 13 RUSSIAPAT: New full-text patent database on STN
NEWS 21 JUN 13 FRFULL enhanced with patent drawing images
NEWS 22 JUN 27 MARPAT displays enhanced with expanded G-group definitions
and text labels
NEWS 23 JUL 01 MEDICONF removed from STN
NEWS 24 JUL 07 STN Patent Forums to be held in July 2005
NEWS 25 JUL 13 SCISEARCH reloaded
NEWS 26 JUL 20 Powerful new interactive analysis and visualization software,
STN AnaVist, now available

NEWS EXPRESS JUNE 13 CURRENT WINDOWS VERSION IS V8.0, CURRENT
MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP),
AND CURRENT DISCOVER FILE IS DATED 13 JUNE 2005

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FILE 'HOME' ENTERED AT 17:59:30 ON 23 JUL 2005

=> file reg

COST IN U.S. DOLLARS

SINCE FILE
ENTRY

TOTAL
SESSION

10/773,002

FULL ESTIMATED COST

0.21

0.21

FILE 'REGISTRY' ENTERED AT 17:59:42 ON 23 JUL 2005
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STRUCTURE FILE UPDATES: 22 JUL 2005 HIGHEST RN 856698-04-9
DICTIONARY FILE UPDATES: 22 JUL 2005 HIGHEST RN 856698-04-9

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TSCA INFORMATION NOW CURRENT THROUGH JANUARY 18, 2005

Please note that search-term pricing does apply when
conducting SmartSELECT searches.

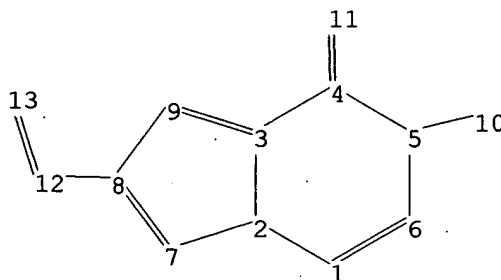
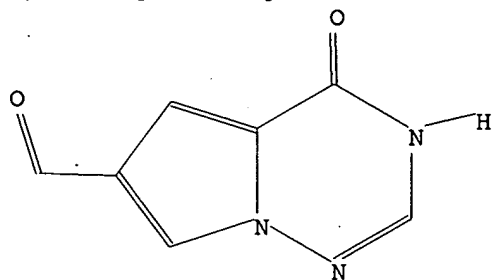
*
* The CA roles and document type information have been removed from *
* the IDE default display format and the ED field has been added, *
* effective March 20, 2005. A new display format, IDERL, is now *
* available and contains the CA role and document type information. *
*

Structure search iteration limits have been increased. See HELP SLIMITS
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Experimental and calculated property data are now available. For more
information enter HELP PROP at an arrow prompt in the file or refer
to the file summary sheet on the web at:
<http://www.cas.org/ONLINE/DBSS/registryss.html>

=>

Uploading C:\Program Files\Stnexp\Queries\10773002B2.str



chain nodes :
10 11 12 13
ring nodes :
1 2 3 4 5 6 7 8 9
chain bonds :
4-11 5-10 8-12 12-13
ring bonds :
1-2 1-6 2-3 2-7 3-4 3-9 4-5 5-6 7-8 8-9
exact/norm bonds :
1-2 1-6 2-3 2-7 3-4 3-9 4-5 4-11 5-6 7-8 8-9 12-13
exact bonds :

10/773,002

5-10 8-12

Match level :

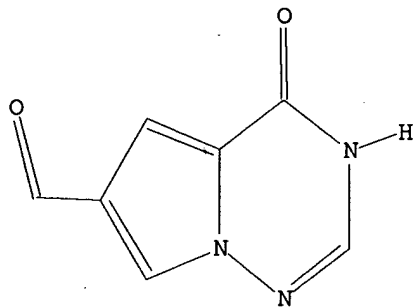
1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom 10:CLASS
11:CLASS 12:CLASS 13:CLASS

L1 STRUCTURE UPLOADED

=> d l1

L1 HAS NO ANSWERS

L1 STR



Structure attributes must be viewed using STN Express query preparation.

=> s l1 sss sam

SAMPLE SEARCH INITIATED 18:00:04 FILE 'REGISTRY'

SAMPLE SCREEN SEARCH COMPLETED - 14 TO ITERATE

100.0% PROCESSED 14 ITERATIONS

0 ANSWERS

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**
BATCH **COMPLETE**

PROJECTED ITERATIONS: 56 TO 504

PROJECTED ANSWERS: 0 TO 0

L2 0 SEA SSS SAM L1

=> s l1 sss ful

FULL SEARCH INITIATED 18:00:14 FILE 'REGISTRY'

FULL SCREEN SEARCH COMPLETED - 487 TO ITERATE

100.0% PROCESSED 487 ITERATIONS

23 ANSWERS

SEARCH TIME: 00.00.01

L3 23 SEA SSS FUL L1

=> file caplus

COST IN U.S. DOLLARS

SINCE FILE

TOTAL

ENTRY

SESSION

FULL ESTIMATED COST

161.33

161.54

FILE 'CAPLUS' ENTERED AT 18:00:20 ON 23 JUL 2005

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FILE COVERS 1907 - 23 Jul 2005 VOL 143 ISS 5
 FILE LAST UPDATED: 22 Jul 2005 (20050722/ED)

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This file contains CAS Registry Numbers for easy and accurate substance identification.

=> s 13

L4 15 L3

=> d 14 1-15 bib hitstr

L4 ANSWER 1 OF 15 CAPLUS COPYRIGHT 2005 ACS on STN

AN 2005:567104 CAPLUS

TI A preparation of (indazolylamino)pyrrolo[2,1-f][1,2,4]triazine derivatives, useful as antiproliferative agents

IN Swaminathan, Shankar; Gavai, Ashvinikumar V.; Fan, Junying; Patel, Bharat P.; Norris, Derek J.; Corbett, Richard Michael; Zheng, Bin

PA Bristol-Myers Squibb Company, USA

SO PCT Int. Appl., 51 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2005058245	A2	20050630	WO 2004-US41920	20041210
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	RW:				
	BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				

PRAI US 2003-529347P P 20031212

US 2004-8719 A 20041209

IT 427878-70-4

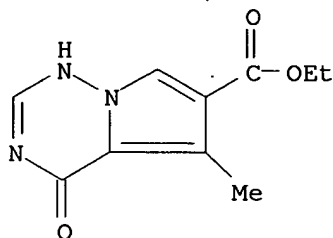
RL: RCT (Reactant); RACT (Reactant or reagent)

(preparation of (indazolylamino)pyrrolo[2,1-f][1,2,4]triazine derivs. useful as antiproliferative agents)

RN 427878-70-4 CAPLUS

10/773,002

CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-oxo-, ethyl ester (9CI) (CA INDEX NAME)



L4 ANSWER 2 OF 15 CAPLUS COPYRIGHT 2005 ACS on STN

AN 2005:160831 CAPLUS

DN 142:261564

TI Heteroaryl-substituted pyrrolo-triazine compounds useful as kinase inhibitors, particularly p38 kinases, and their preparation, pharmaceutical compositions, and use

IN Leftheris, Katerina; Wroblewski, Stephen T.; Dyckman, Alaric J.

PA USA

SO U.S. Pat. Appl. Publ., 29 pp., Cont.-in-part of U.S. Ser. No. 420,399.
CODEN: USXXCO

DT Patent

LA English

FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 2005043306	A1	20050224	US 2003-678388	20031003
	US 2004082582	A1	20040429	US 2003-420399	20030422
	WO 2005037838	A1	20050428	WO 2004-US30829	20040921
	W:				
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	RW:				
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PRAI US 2002-374938P P 20020423

US 2003-420399 A2 20030422

US 2003-678388 A 20031003

OS MARPAT 142:261564

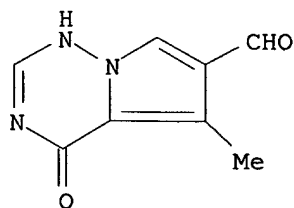
IT 621685-55-0P 621685-56-1P 621685-58-3P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(intermediate; preparation of heteroaryl pyrrolotriazine compds. as p38 kinase inhibitors)

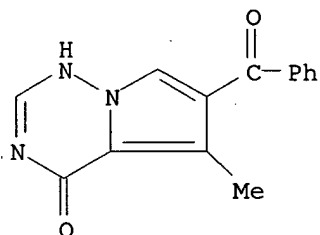
RN 621685-55-0 CAPLUS

CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxaldehyde, 1,4-dihydro-5-methyl-4-oxo- (9CI) (CA INDEX NAME)



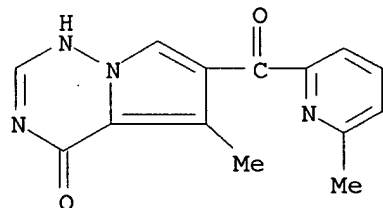
RN 621685-56-1 CAPLUS

CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one, 6-benzoyl-5-methyl- (9CI) (CA INDEX NAME)



RN 621685-58-3 CAPLUS

CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one, 5-methyl-6-[(6-methyl-2-pyridinyl)carbonyl]- (9CI) (CA INDEX NAME)



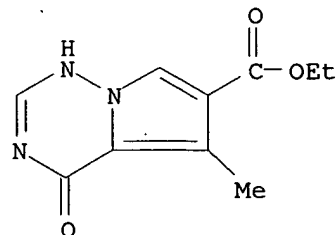
IT 427878-70-4

RL: RCT (Reactant); RACT (Reactant or reagent)

(preparation of heteroaryl pyrrolotriazine compds. as p38 kinase inhibitors)

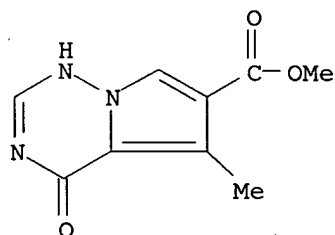
RN 427878-70-4 CAPLUS

CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-oxo-, ethyl ester (9CI) (CA INDEX NAME)

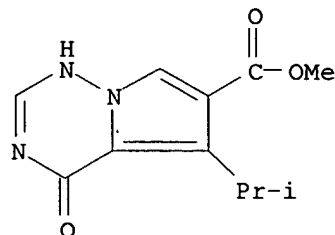


10/773,002

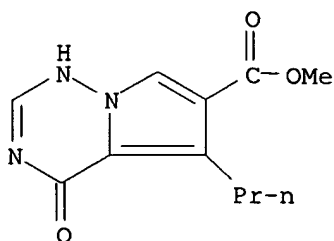
L4 ANSWER 3 OF 15 CAPLUS COPYRIGHT 2005 ACS on STN
AN 2005:130304 CAPLUS
DN 142:392380
TI Synthesis and SAR of 4-(3-hydroxyphenylamino)pyrrolo[2,1-f][1,2,4]triazine based VEGFR-2 kinase inhibitors
AU Borzilleri, Robert M.; Cai, Zhen-Wei; Ellis, Christopher; Fargnoli, Joseph; Fura, Abera; Gerhardt, Tracy; Goyal, Bindu; Hunt, John T.; Mortillo, Steven; Qian, Ligang; Tokarski, John; Vyas, Viral; Wautlet, Barri; Zheng, Xioping; Bhide, Rajeev S.
CS Departments of Oncology Chemistry, Bristol-Myers Squibb Pharmaceutical Research Institute, Princeton, NJ, 08543-4000, USA
SO Bioorganic & Medicinal Chemistry Letters (2005), 15(5), 1429-1433
CODEN: BMCLE8; ISSN: 0960-894X
PB Elsevier B.V.
DT Journal
LA English
IT 310431-29-9P 651744-51-3P 850085-64-2P
850085-66-4P 850085-97-1P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(synthesis and SAR of 4-(3-hydroxyphenylamino)pyrrolo[2,1-f][1,2,4]triazine based VEGFR-2 kinase inhibitors)
RN 310431-29-9 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-oxo-, methyl ester (9CI) (CA INDEX NAME)



RN 651744-51-3 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-(1-methylethyl)-4-oxo-, methyl ester (9CI) (CA INDEX NAME)

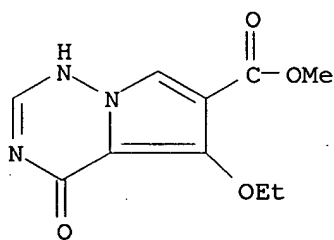


RN 850085-64-2 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-4-oxo-5-propyl-, methyl ester (9CI) (CA INDEX NAME)



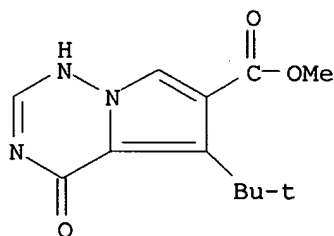
RN 850085-66-4 CAPLUS

CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 5-ethoxy-1,4-dihydro-4-oxo-, methyl ester (9CI) (CA INDEX NAME)



RN 850085-97-1 CAPLUS

CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 5-(1,1-dimethylethyl)-1,4-dihydro-4-oxo-, methyl ester (9CI) (CA INDEX NAME)



RE.CNT 13 THERE ARE 13 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 4 OF 15 CAPLUS COPYRIGHT 2005 ACS on STN

AN 2004:654775 CAPLUS

DN 141:190807

TI Process for preparing pyrrolotriazine kinase inhibitors

IN Chen, Bang-Chi; Zhao, Rulin; Sundeen, Joseph Edward; Leftheris, Katerina; Hynes, John; Wroblewski, Stephen T.

PA USA

SO U.S. Pat. Appl. Publ., 20 pp.

CODEN: USXXCO

DT Patent

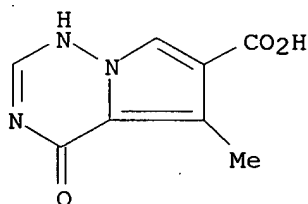
LA English

FAN.CNT 1

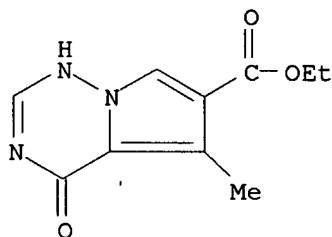
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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10/773,002

PI US 2004157846 A1 20040812 US 2004-773002 20040205
WO 2004072030 A2 20040826 WO 2004-US3223 20040205
WO 2004072030 A3 20041028
W: AE, AE, AG, AL, AL, AM, AM, AM, AT, AT, AU, AZ, AZ, BA, BB, BG,
BG, BR, BR, BW, BY, BY, BZ, BZ, CA, CH, CN, CN, CO, CO, CR, CR,
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RW: BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE,
BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU,
MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN,
GQ, GW, ML, MR, NE, SN, TD, TG, BF, BJ, CF, CG, CI, CM, GA, GN,
GQ, GW, ML, MR, NE, SN, TD, TG
PRAI US 2003-445224P P 20030205
OS CASREACT 141:190807; MARPAT 141:190807
IT 310435-15-5P 427878-70-4P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
(Reactant or reagent)
(process for preparing pyrrolotriazine p38 kinase inhibitors)
RN 310435-15-5 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-
oxo- (9CI) (CA INDEX NAME)



RN 427878-70-4 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-
oxo-, ethyl ester (9CI) (CA INDEX NAME)



L4 ANSWER 5 OF 15 CAPLUS COPYRIGHT 2005 ACS on STN
AN 2004:531311 CAPLUS
DN 141:89122
TI Preparation of C-6 modified indazolyl pyrrolotriazines as
antiproliferative agents
IN Vite, Gregory D.; Gavai, Ashvinikumar V.; Fink, Brian E.; Mastalerz,
Harold; Kadow, John F.
PA Bristol-Myers Squibb Company, USA
SO PCT Int. Appl., 81 pp.

10/773,002

CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

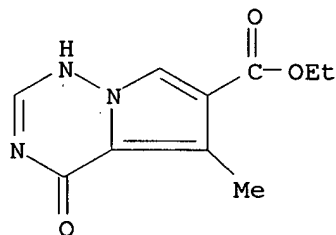
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PI	WO 2004054514	A2	20040701	WO 2003-US39542	20031212
	WO 2004054514	A3	20041007		
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	RW: BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
	US 2004142931	A1	20040722	US 2003-736476	20031215
	US 6916815	B2	20050712		
PRAI	US 2002-433190P	P	20021213		
OS	MARPAT 141:89122				
IT	427878-70-4 714971-30-9				

RL: RCT (Reactant); RACT (Reactant or reagent)

(preparation of C-6 modified indazolyl pyrrolotriazines as antiproliferative agents)

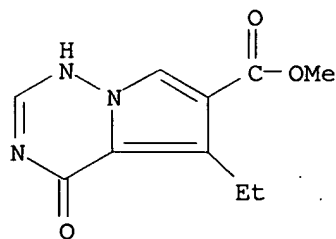
RN 427878-70-4 CAPLUS

CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-oxo-, ethyl ester (9CI) (CA INDEX NAME)



RN 714971-30-9 CAPLUS

CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 5-ethyl-1,4-dihydro-4-oxo-, methyl ester (9CI) (CA INDEX NAME)



L4 ANSWER 6 OF 15 CAPLUS COPYRIGHT 2005 ACS on STN
AN 2004:120859 CAPLUS

10/773,002

DN 140:181471
TI Preparation of pyrrolotriazines as tyrosine kinase activity inhibitors of growth factor receptors for the treatment of cancer
IN Bhide, Rajeev S.; Borzilleri, Robert M.
PA Bristol-Myers Squibb Company, USA
SO PCT Int. Appl., 71 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2004013145	A1	20040212	WO 2003-US24273	20030804
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	EP 1543009	A1	20050622	EP 2003-767116	20030804
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	WO 2003-US24273	W	20030804		

OS MARPAT 140:181471

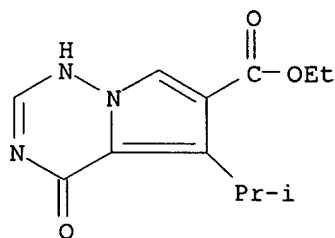
IT 651744-40-0P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(drug candidate; preparation of pyrrolotriazines as tyrosine kinase activity inhibitors of growth factor receptors)

RN 651744-40-0 CAPLUS

CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-(1-methylethyl)-4-oxo-, ethyl ester (9CI) (CA INDEX NAME)



IT 658085-53-1P 658085-61-1P 658085-62-2P

658085-63-3P 658085-64-4P 658085-69-9P

658085-71-3P, 4-Hydroxy-5-(1-methylethyl)pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid(2-oxopropyl)amide

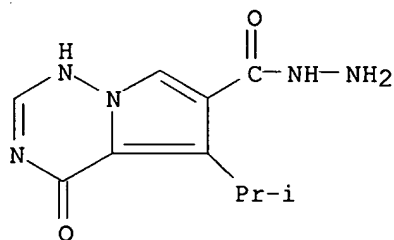
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(intermediate; preparation of pyrrolotriazines as tyrosine kinase activity inhibitors of growth factor receptors)

RN 658085-53-1 CAPLUS

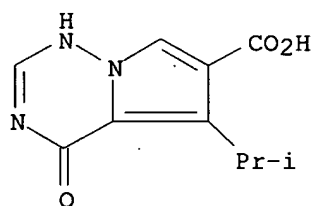
10/773,002

CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-(1-methylethyl)-4-oxo-, hydrazide (9CI) (CA INDEX NAME)



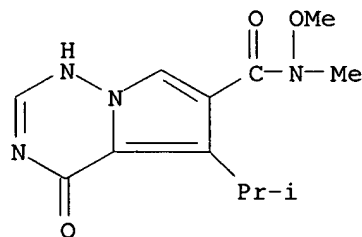
RN 658085-61-1 CAPLUS

CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-(1-methylethyl)-4-oxo- (9CI) (CA INDEX NAME)



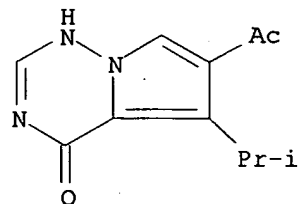
RN 658085-62-2 CAPLUS

CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxamide, 1,4-dihydro-N-methoxy-N-methyl-5-(1-methylethyl)-4-oxo- (9CI) (CA INDEX NAME)



RN 658085-63-3 CAPLUS

CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one, 6-acetyl-5-(1-methylethyl)- (9CI) (CA INDEX NAME)

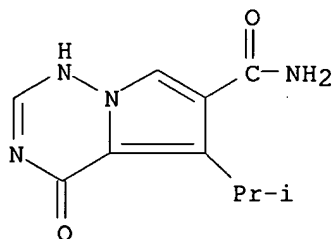


RN 658085-64-4 CAPLUS

CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxamide, 1,4-dihydro-5-(1-methylethyl)-

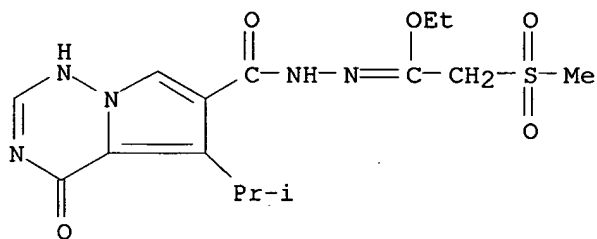
10/773,002

4-oxo- (9CI) (CA INDEX NAME)



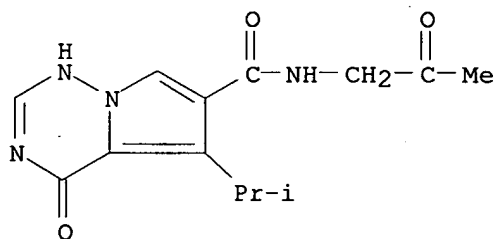
RN 658085-69-9 CAPLUS

CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-(1-methylethyl)-4-oxo-, [1-ethoxy-2-(methylsulfonyl)ethylidene]hydrazide (9CI) (CA INDEX NAME)



RN 658085-71-3 CAPLUS

CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxamide, 1,4-dihydro-5-(1-methylethyl)-4-oxo-N-(2-oxopropyl)- (9CI) (CA INDEX NAME)



L4 ANSWER 7 OF 15 CAPLUS COPYRIGHT 2005 ACS on STN

AN 2004:80847 CAPLUS

DN 140:124558

TI Pyrrolotriazine inhibitors of kinases for use in treatment of diseases associated with growth factor receptor signal transduction

IN Bhide, Rajeev; Cai, Zhen-wei; Qian, Ligang; Barbosa, Stephanie

PA Bristol-Myers Squibb Company, USA

SO PCT Int. Appl., 84 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 3

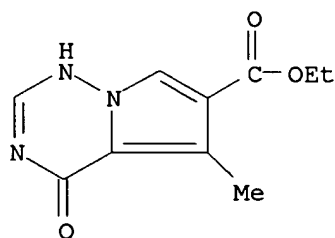
PATENT NO.

KIND DATE

APPLICATION NO.

DATE

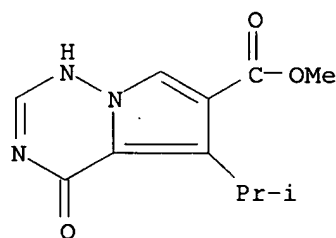
PI WO 2004009784 A2 20040129 WO 2003-US22826 20030718
 WO 2004009784 A3 20040422
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 GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,
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 PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN,
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 BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
 CA 2492804 AA 20040129 CA 2003-2492804 20030718
 US 2004063707 A1 20040401 US 2003-622593 20030718
 US 2004072832 A1 20040415 US 2003-623171 20030718
 US 6869952 B2 20050322
 EP 1534290 A2 20050601 EP 2003-765881 20030718
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 IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK
 US 2005124621 A1 20050609 US 2005-35248 20050113
 PRAI US 2002-397256P P 20020719
 US 2003-447213P P 20030213
 US 2003-623171 A1 20030718
 WO 2003-US22826 W 20030718
 OS MARPAT 140:124558
 IT 427878-70-4
 RL: RCT (Reactant); RACT (Reactant or reagent)
 (pyrrolotriazine inhibitors of kinases for use in treatment of diseases
 associated with growth factor receptor signal transduction)
 RN 427878-70-4 CAPLUS
 CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-
 oxo-, ethyl ester (9CI) (CA INDEX NAME)



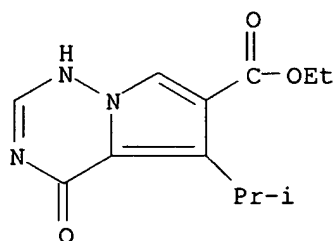
L4 ANSWER 8 OF 15 CAPLUS COPYRIGHT 2005 ACS on STN
 AN 2004:80698 CAPLUS
 DN 140:146173
 TI Preparation of pyrrolotriazines as selective VEGFR-2 and FGFR-1 kinase
 inhibitors for treatment of proliferative diseases
 IN Bhide, Rajeev; Ruel, Rejean; Thibeault, Carl; L'heureux, Alexandre
 PA Bristol-Myers Squibb Company, USA
 SO PCT Int. Appl., 66 pp.
 CODEN: PIXXD2
 DT Patent
 LA English
 FAN.CNT 3

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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PI WO 2004009601 A1 20040129 WO 2003-US22554 20030718
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RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
CA 2492665 AA 20040129 CA 2003-2492665 20030718
US 2004063707 A1 20040401 US 2003-622593 20030718
US 2004072832 A1 20040415 US 2003-623171 20030718
US 6869952 B2 20050322
EP 1539763 A1 20050615 EP 2003-765754 20030718
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK
US 2005124621 A1 20050609 US 2005-35248 20050113
PRAI US 2002-397256P P 20020719
US 2003-447213P P 20030213
US 2003-623171 A1 20030718
WO 2003-US22554 W 20030718
OS MARPAT 140:146173
IT **651744-51-3**
RL: RCT (Reactant); RACT (Reactant or reagent)
(preparation of pyrrolotriazines as selective VEGFR-2 and FGFR-1 kinase inhibitors for treatment of proliferative diseases)
RN 651744-51-3 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-(1-methylethyl)-4-oxo-, methyl ester (9CI) (CA INDEX NAME)



IT **651744-40-0P**
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation of pyrrolotriazines as selective VEGFR-2 and FGFR-1 kinase inhibitors for treatment of proliferative diseases)
RN 651744-40-0 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-(1-methylethyl)-4-oxo-, ethyl ester (9CI) (CA INDEX NAME)



RE.CNT 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 9 OF 15 CAPLUS COPYRIGHT 2005 ACS on STN

AN 2004:80644 CAPLUS

DN 140:146018

TI Process for preparation of indolyloxypyrrolotriazines and their use as drugs.

IN Bhide, Rajeev; Fan, Junying; Parlanti, Luca; Barbosa, Stephanie; Qian, Ligang; Cai, Zhen-wei; Gibson, Francis S.

PA Bristol-Myers Squibb Company, USA

SO PCT Int. Appl., 48 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 3

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2004009542	A2	20040129	WO 2003-US22755	20030721
	WO 2004009542	A3	20040513		
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	US 2004077858	A1	20040422	US 2003-622280	20030718
	CA 2492861	AA	20040129	CA 2003-2492861	20030721
	EP 1554281	A2	20050720	EP 2003-765846	20030721
	R:	AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK			
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PRAI	US 2002-397256P	P	20020719		
	US 2003-447213P	P	20030213		
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	WO 2003-US22755	W	20030721		

OS MARPAT 140:146018

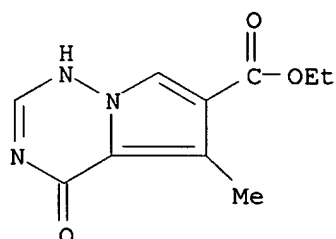
IT 427878-70-4

RL: RCT (Reactant); RACT (Reactant or reagent)

(process for preparation of indolyloxypyrrolotriazines and their use as drugs)

RN 427878-70-4 CAPLUS

CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-oxo-, ethyl ester (9CI) (CA INDEX NAME)



L4 ANSWER 10 OF 15 CAPLUS COPYRIGHT 2005 ACS on STN

AN 2003:875265 CAPLUS

DN 139:364963

TI Aryl ketone pyrrolo-triazine compounds useful as kinase inhibitors, particularly p38 kinases, and their preparation, pharmaceutical compositions, and use

IN Dyckman, Alaric; Leftheris, Katerina; Hynes, John

PA Bristol-Myers Squibb Company, USA

SO PCT Int. Appl., 45 pp.

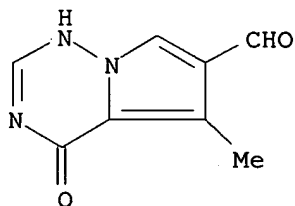
CODEN: PIXXD2

DT Patent

LA English

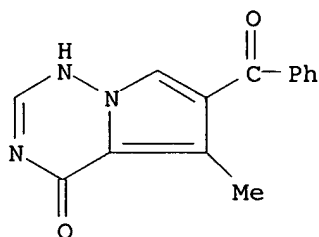
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PI	WO 2003091229	A1	20031106	WO 2003-US12420	20030418
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	EP 1503996	A1	20050209	EP 2003-718493	20030418
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
	US 2003232831	A1	20031218	US 2003-420445	20030422
PRAI	US 2002-374907P	P	20020423		
	WO 2003-US12420	W	20030418		
OS	MARPAT 139:364963				
IT	621685-55-0P 621685-56-1P 621685-58-3P				
	RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)				
	(intermediate; preparation of aryl ketone pyrrolotriazine compds. as p38 kinase inhibitors)				
RN	621685-55-0 CAPLUS				
CN	Pyrrolo[2,1-f][1,2,4]triazine-6-carboxaldehyde, 1,4-dihydro-5-methyl-4-oxo-(9CI) (CA INDEX NAME)				



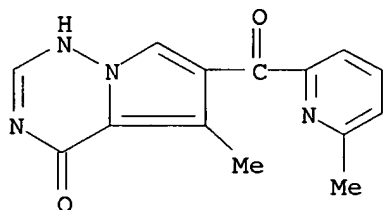
RN 621685-56-1 CAPLUS

CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one, 6-benzoyl-5-methyl- (9CI) (CA INDEX NAME)



RN 621685-58-3 CAPLUS

CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one, 5-methyl-6-[(6-methyl-2-pyridinyl)carbonyl]- (9CI) (CA INDEX NAME)

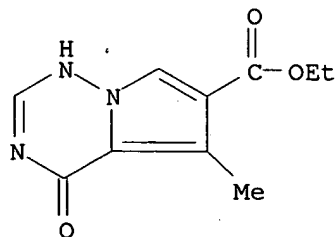


IT 427878-70-4

RL: RCT (Reactant); RACT (Reactant or reagent)
(starting material; preparation of aryl ketone pyrrolotriazine compds. as p38 kinase inhibitors)

RN 427878-70-4 CAPLUS

CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-oxo-, ethyl ester (9CI) (CA INDEX NAME)

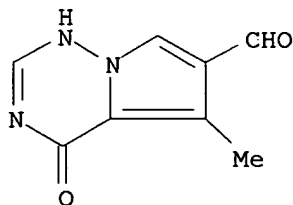


10/773,002

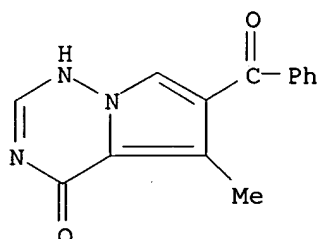
RE.CNT 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 11 OF 15 CAPLUS COPYRIGHT 2005 ACS on STN
AN 2003:875173 CAPLUS
DN 139:381511
TI Pyrrolotriazine aniline compounds useful as kinase inhibitors,
particularly p38 kinases, and their preparation, pharmaceutical
compositions, and use as antiinflammatory agents
IN Dyckman, Alaric; Hynes, John; Leftheris, Katherina; Liu, Chunjian;
Wroblewski, Stephen T.
PA Bristol-Myers Squibb Company, USA
SO PCT Int. Appl., 158 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2003090912	A1	20031106	WO 2003-US12426	20030415
	WO 2003090912	C2	20040108		
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	EP 1497019	A1	20050119	EP 2003-724157	20030415
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PRAI	US 2002-374938P	P	20020423		
	WO 2003-US12426	W	20030415		
OS	MARPAT 139:381511				
IT	621685-55-0P 621685-56-1P 621685-58-3P				
	RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)				
	(intermediate; preparation of pyrrolotriazine aniline compds. as p38 kinase inhibitors)				
RN	621685-55-0 CAPLUS				
CN	Pyrrolo[2,1-f][1,2,4]triazine-6-carboxaldehyde, 1,4-dihydro-5-methyl-4-oxo- (9CI) (CA INDEX NAME)				

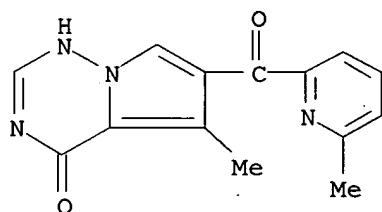


RN 621685-56-1 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one, 6-benzoyl-5-methyl- (9CI) (CA INDEX NAME)



RN 621685-58-3 CAPLUS

CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one, 5-methyl-6-[(6-methyl-2-pyridinyl)carbonyl]- (9CI) (CA INDEX NAME)



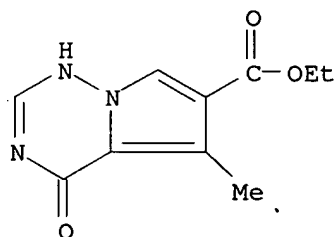
IT 427878-70-4

RL: RCT (Reactant); RACT (Reactant or reagent)

(starting material; preparation of pyrrolo-triazine aniline compds. as p38 kinase inhibitors)

RN 427878-70-4 CAPLUS

CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-oxo-, ethyl ester (9CI) (CA INDEX NAME)



RE.CNT 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 12 OF 15 CAPLUS COPYRIGHT 2005 ACS on STN

AN 2003:777390 CAPLUS

DN 139:292275

TI Methods for the preparation of pyrrolo-triazine compounds useful as kinase inhibitors

IN Godfrey, Jollie Duaine; Hynes, John; Dyckman, Alaric J.; Leftheris, Katerina; Shi, Zhongping; Wroblewski, Stephen T.; Doubleday, Wendel William; Grosso, John A.

PA Bristol-Myers Squibb Company, USA

SO U.S. Pat. Appl. Publ., 36 pp., Cont.-in-part of U.S. Ser. No. 36,293.

10/773,002

CODEN: USXXCO
DT Patent
LA English
FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 2003186982	A1	20031002	US 2002-289010	20021106
	US 6867300	B2	20050315		
	US 2003069244	A1	20030410	US 2001-36293	20011107
	US 6670357	B2	20031230		
	ZA 2003003786	A	20040816	ZA 2003-3786	20030515
	US 2004229877	A1	20041118	US 2003-696178	20031029
	WO 2004043912	A2	20040527	WO 2003-US35220	20031103
	WO 2004043912	A3	20040701		

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW

RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

	US 2005107462	A1	20050519	US 2004-19788	20041222
PRAI	US 2000-249877P	P	20001117		
	US 2001-310561P	P	20010807		
	US 2001-36293	A2	20011107		
	US 2002-289010	A	20021106		

OS MARPAT 139:292275

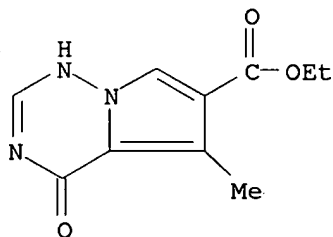
IT **427878-70-4P**

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(preparation of pyrrolotriazine derivative as kinase inhibitor)

RN 427878-70-4 CAPLUS

CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-oxo-, ethyl ester (9CI) (CA INDEX NAME)



L4 ANSWER 13 OF 15 CAPLUS COPYRIGHT 2005 ACS on STN

AN 2003:396849 CAPLUS

DN 138:401758

TI Preparation of 5-substituted N-(1H-indazol-5-yl)pyrrolo[2,1-f][1,2,4]triazin-4-amines as antiproliferative agents

IN Mastalerz, Harold; Zhang, Guifen; Tarrant, James G.; Vite, Gregory D.

PA Bristol-Myers Squibb Company, USA

SO PCT Int. Appl., 74 pp.

CODEN: PIXXD2

DT Patent

LA English
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2003042172	A2	20030522	WO 2002-US36528	20021112
	WO 2003042172	A3	20040129		
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
	CA 2467068	AA	20030522	CA 2002-2467068	20021112
	EP 1446401	A2	20040818	EP 2002-793930	20021112
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, SK				
	BR 2002014112	A	20040914	BR 2002-14112	20021112
	NZ 533034	A	20041126	NZ 2002-533034	20021112
	JP 2005509030	T2	20050407	JP 2003-544009	20021112
	US 2003186983	A1	20031002	US 2002-294281	20021114
	US 6908916	B2	20050621		
PRAI	US 2001-333014P	P	20011114		
	WO 2002-US36528	W	20021112		

OS MARPAT 138:401758

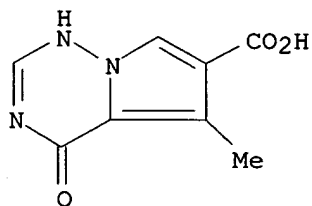
IT **310435-15-5P**, 5-Methyl-4-oxo-3,4-dihydropyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(intermediate; preparation of N-(indazolyl)pyrrolotriazinamines as tyrosine kinase inhibitors for treatment of proliferative disorders and other diseases associated with signal transduction pathways)

RN 310435-15-5 CAPLUS

CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-oxo- (9CI) (CA INDEX NAME)



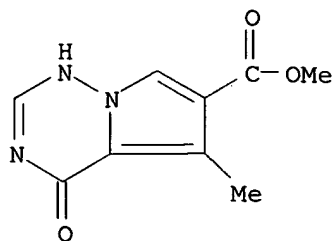
IT **310431-29-9**, 5-Methyl-4-oxo-3,4-dihydropyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid methyl ester

RL: RCT (Reactant); RACT (Reactant or reagent)

(preparation of N-(indazolyl)pyrrolotriazinamines as tyrosine kinase inhibitors for treatment of proliferative disorders and other diseases associated with signal transduction pathways)

RN 310431-29-9 CAPLUS

CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-oxo-, methyl ester (9CI) (CA INDEX NAME)

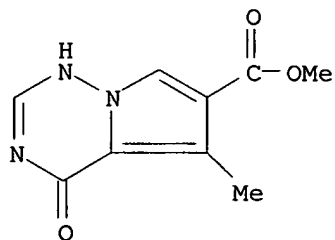


L4 ANSWER 14 OF 15 CAPLUS COPYRIGHT 2005 ACS on STN
 AN 2002:391720 CAPLUS
 DN 136:386144
 TI Preparation of pyrrolo[2,1-f][1,2,4]triazine carboxylic acid derivatives
 for use in treating p38 kinase-associated conditions
 IN Leftheris, Katerina; Barrish, Joel; Hynes, John; Wroblewski, Stephen T.
 PA Bristol-Myers Squibb Company, USA
 SO PCT Int. Appl., 108 pp.
 CODEN: PIXXD2
 DT Patent
 LA English
 FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2002040486	A2	20020523	WO 2001-US49982	20011107
	WO 2002040486	A3	20030912		
	W:				
	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,				
	CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,				
	GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,				
	LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PH, PL,				
	PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG,				
	US, UZ, VN, YU, ZA, ZW				
	RW:				
	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AM, AZ, BY, KG,				
	KZ, MD, RU, TJ, TM, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR,				
	IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN,				
	GQ, GW, ML, MR, NE, SN, TD, TG				
	CA 2429628	AA	20020523	CA 2001-2429628	20011107
	AU 2002032760	A5	20020527	AU 2002-32760	20011107
	EE 200300227	A	20031015	EE 2003-227	20011107
	EP 1363910	A2	20031126	EP 2001-992298	20011107
	R:				
	AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,				
	IE, SI, LT, LV, FI, RO, MK, CY, AL, TR				
	JP 2004522713	T2	20040729	JP 2002-543494	20011107
	BG 107750	A	20040130	BG 2003-107750	20030421
	ZA 2003003786	A	20040816	ZA 2003-3786	20030515
	NO 2003002229	A	20030716	NO 2003-2229	20030516
PRAI	US 2000-249877P	P	20001117		
	US 2001-310561P	P	20010807		
	WO 2001-US49982	W	20011107		
OS	MARPAT 136:386144				
IT	310431-29-9P 310435-15-5P 310443-54-0P,				
	4-Hydroxy-5-methoxypyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid ethyl				
	ester 427878-70-4P				
	RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT				
	(Reactant or reagent)				
	(intermediate; preparation of pyrrolo[2,1-f][1,2,4]triazine carboxylic acid				
	derivs. for use in treating p38 kinase-associated conditions)				
RN	310431-29-9 CAPLUS				

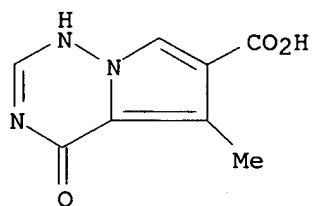
10/773,002

CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-oxo-, methyl ester (9CI) (CA INDEX NAME)



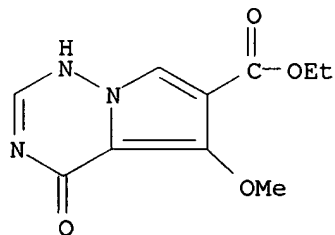
RN 310435-15-5 CAPLUS

CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-oxo- (9CI) (CA INDEX NAME)



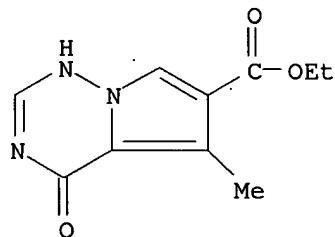
RN 310443-54-0 CAPLUS

CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methoxy-4-oxo-, ethyl ester (9CI) (CA INDEX NAME)



RN 427878-70-4 CAPLUS

CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-oxo-, ethyl ester (9CI) (CA INDEX NAME)



L4 ANSWER 15 OF 15 CAPLUS COPYRIGHT 2005 ACS on STN
 AN 2000:841986 CAPLUS
 DN 134:17506
 TI Preparation of pyrrolotriazines as kinases inhibitors for treating
 inflammation, cancer, and proliferative diseases
 IN Hunt, John T.; Bhide, Rajeev S.; Borzilleri, Robert M.; Qian, Ligang
 PA Bristol-Myers Squibb Company, USA
 SO PCT Int. Appl., 130 pp.
 CODEN: PIXXD2
 DT Patent
 LA English
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2000071129	A1	20001130	WO 2000-US13420	20000516
	W:	AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			
	RW:	GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG			
	CA 2373990	AA	20001130	CA 2000-2373990	20000516
	EP 1183033	A1	20020306	EP 2000-930761	20000516
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	BR 2000010482	A	20020423	BR 2000-10482	20000516
	JP 2003500359	T2	20030107	JP 2000-619433	20000516
	NZ 516292	A	20040130	NZ 2000-516292	20000516
	AU 770377	B2	20040219	AU 2000-48524	20000516
	TR 200103352	T2	20050321	TR 2001-200103352	20000516
	NO 2001005650	A	20011120	NO 2001-5650	20011120
	ZA 2001009577	A	20030220	ZA 2001-9577	20011120
PRAI	US 1999-135265P	P	19990521		
	US 2000-193727P	P	20000331		
	WO 2000-US13420	W	20000516		

OS MARPAT 134:17506

IT 310431-16-4P 310431-29-9P 310435-15-5P

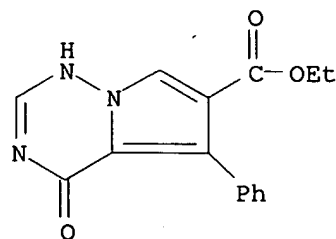
310436-48-7P 310436-60-3P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(preparation of pyrrolotriazines as kinases inhibitors useful in treating inflammation, cancer, and proliferative diseases)

RN 310431-16-4 CAPLUS

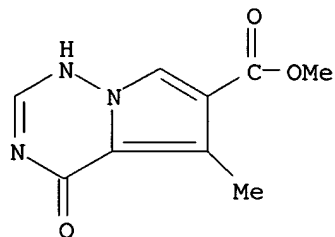
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-4-oxo-5-phenyl-, ethyl ester (9CI) (CA INDEX NAME)



10/773,002

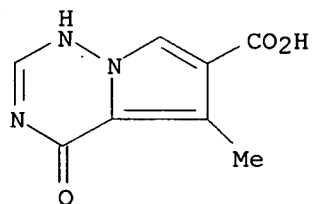
RN 310431-29-9 CAPLUS

CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-oxo-, methyl ester (9CI) (CA INDEX NAME)



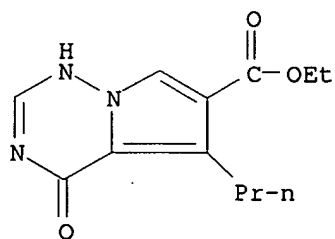
RN 310435-15-5 CAPLUS

CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-oxo- (9CI) (CA INDEX NAME)



RN 310436-48-7 CAPLUS

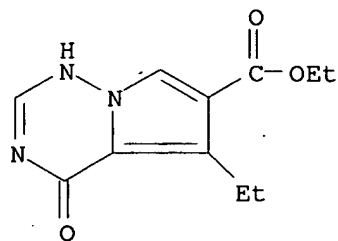
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-4-oxo-5-propyl-, ethyl ester (9CI) (CA INDEX NAME)



RN 310436-60-3 CAPLUS

CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 5-ethyl-1,4-dihydro-4-oxo-, ethyl ester (9CI) (CA INDEX NAME)

10/773,002



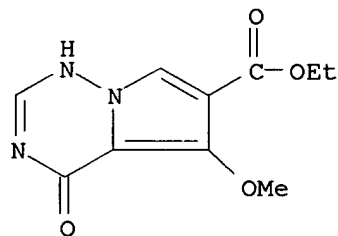
IT 310443-54-0P

RL: SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of pyrrolotriazines as kinases inhibitors useful in treating inflammation, cancer, and proliferative diseases)

RN 310443-54-0 CAPLUS

CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methoxy-4-oxo-, ethyl ester (9CI) (CA INDEX NAME)



RE.CNT 8 THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

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COST IN U.S. DOLLARS

SINCE FILE

TOTAL

ENTRY

SESSION

FULL ESTIMATED COST

51.30

212.84

STN INTERNATIONAL LOGOFF AT 18:00:44 ON 23 JUL 2005

Ref #	Hits	Search Query	DBs	Default Operator	Plurals	Time Stamp
L1	344	(544/183).CCLS.	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/07/23 19:16
L2	40	Bang-Chi.inv. and Chen.inv.	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/07/23 19:16
L3	28	Rulin.inv. and Zhao.inv.	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/07/23 19:17
L4	79	Joseph.inv. and Sundeen.inv.	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/07/23 19:17
L5	57	katerina.inv. and Leftheris.inv.	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/07/23 19:18
L6	68	John.inv. and Hynes.inv.	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/07/23 19:18
L7	0	Stephen.inv. and Wrobieski.inv.	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/07/23 19:19
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Ref #	Hits	Search Query	DBs	Default Operator	Plurals	Time Stamp
L1	344	(544/183).CCLS.	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/07/23 19:16
L2	40	Bang-Chi.inv. and Chen.inv.	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/07/23 19:16
L3	28	Rulin.inv. and Zhao.inv.	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/07/23 19:17
L4	79	Joseph.inv. and Sundeen.inv.	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/07/23 19:17
L5	57	katerina.inv. and Leftheris.inv.	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/07/23 19:18
L6	68	John.inv. and Hynes.inv.	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/07/23 19:18
L7	0	Stephen.inv. and Wrobieski.inv.	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/07/23 19:19
L8	25	Stephen.inv. and Wrobleski.inv.	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/07/23 19:19
L9	7	1 and (2 or 3 or 4 or 5 or 6 or 8)	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/07/23 19:20